Briefings for Michal

- PRIA/registration overview 2/23rd (complete)
- Dicamba 2/18th (complete)
- Sulfoxaflor 3/4th (complete)
- Pet incidents 3/31st (complete)
- FDA/EPA jurisdiction 4/12th (complete)
- SW2 revocation options 5/3rd (complete)
- Antibiotics 5/17th (complete)

New Active Ingredients (include original PRIA date)

- Tetraniliprole completed!
- Broflanilide completed!
- Picarbutrazox completed!
- Fluxametamide completed!
- Fluindapyr completed!
- Trifludimoxazin completed!
- Rekleml PD being developed. PRIA date 7/19/21 (Original PRIA date 4/19/21) (IVB3). Partial ESA causing delays
- Ipflufenoquin PD being developed. PRIA date 7/30/31 (Original PRIA date 4/19/21) (FB). Working on ESA causing delays
- L-Glufosinate Ammonium Risk Assessment phase. PRIA date 10/12/2021 (FHB). ESA issue. Briefing Ed on ESA and resource considerations in June

Other Chemical Actions

- Dicamba
 - Bayer and BASF requesting feedback on pending pre-mixes

Ex. 5 Deliberative Process (DP)

- USDA conducting EIS (instead of EA) on dicamba resistant corn; issued a 60-day public comment period 5/8th; citing EPA's cancellation order and 2020 decision. Discussed with OPMP. Comment period extended to 6/28. Coordinating with AO's OP and regions; OPP team recommends not sending substantive comments at this point
- 24 (c)s -
 - NC denial 3/15th
 - TN denial $4/9^{th}$
 - GA denial 4/23rd
 - TX Notice of intent to deny NLT 6/18th; denial NLT 6/30th draft NOID ready for OPP-IO
- Antibiotics will need to brief the new team (FB)

- Briefed Michal 5/17th; Received policy feedback on the antibiotic issue; Oxytet cherry PRIA date of June delay the action while we work on policy approach.
- **Inorganic Bromide** R10 meeting with OPP; APHIS meeting with Michal requesting increase in tolerance on alfalfa (Dan R)
- Cyclaniliprole/flonicamid –first residential use; public process; published proposal!
- **Seresto** pet collar with high rate of incidents (death and neuro); press inquiries; 6a2 letter sent; updated desk statement to include factual information on FDA vs. EPA; received information from Elanco; team is developing a schedule
- Aldicarb AgLogic is not interested in voluntary cancellation
- Enlist Extension of the registration (expires end of 2021); Corteva is looking for a decision to inform seed purchase for soybeans; OGC advises an to update ESA finding

Other Actions - AA-Level/OD/High Visibility Items

- FDA/EPA jurisdiction Michal follow up complete; team identified appropriate "model" MOUs and sent talking points; next steps on reaching out?
- COVID-19 Activities Sect 18s
 - SW2 -
 - revocation of existing AR, TX S18s based on science and violations (week of June 23rd)
 - Discussing whether AR or OK want to request revocation; OK is requesting revocation grant the request next week (Ed's signature)
 - Grignard reviews complete; working on authorization letters meeting with

Ex. 5 Deliberative Process (DP)

amendment to TN and GA)

- Kraton completed!
- Ionopure Air risk issues
- Acteev Facemasks
- OIG Audits

Ex. 5 Deliberative Process (DP)

- **PFAS** RD working with Clark on submissions
 - Definition issue working to bring Jeff D. up to speed
 - 6a2 notification? Nothing planned yet
- Inerts supply issue phase 1 completed!; phase 2 working on the draft response without AD
- Inert Fragrance petition (Pat Quinn) 2 petition responses scheduled for OGC review in June (Kerry); other petition responses will follow within existing resource constraints

• **CRP** (Child resistant Packaging) testing – industry has requested flexibilities in testing requirements due to COVID impacts; CPSC (whose regulations we use) have issued similar flexibilities. Team recommends issuing some but not all flexibilities. Next steps- COVID workgroup

International

• Bi-monthly meeting with PMRA 5/11th – antibiotics, Seresto – complete; next meeting July

Business/process/technology

- ISB FEP/21-day backlog:
 - backlog addressed by end of April; RD SWAT team addressing newer receipts
 - Working to complete RD SWAT actions
- eCSF/OPPEL eCSF May 9th soft launch; OPPEL -TBD
- MRL structured data for OPP and share with FDA; FDA is applying for internal grant supports OPP digital transformation; OPP serving as reviewer

Speaking Engagements -

- HCPA mid-year meeting Marietta 5/27th
- MRL workshop Nancy Fitz 5/27th

Ex. 5 Deliberative Process (DP)

CA topics (Karen Morrison):

- Missing environmental hazard statements – RD BC's are generating a response